

Figure 1

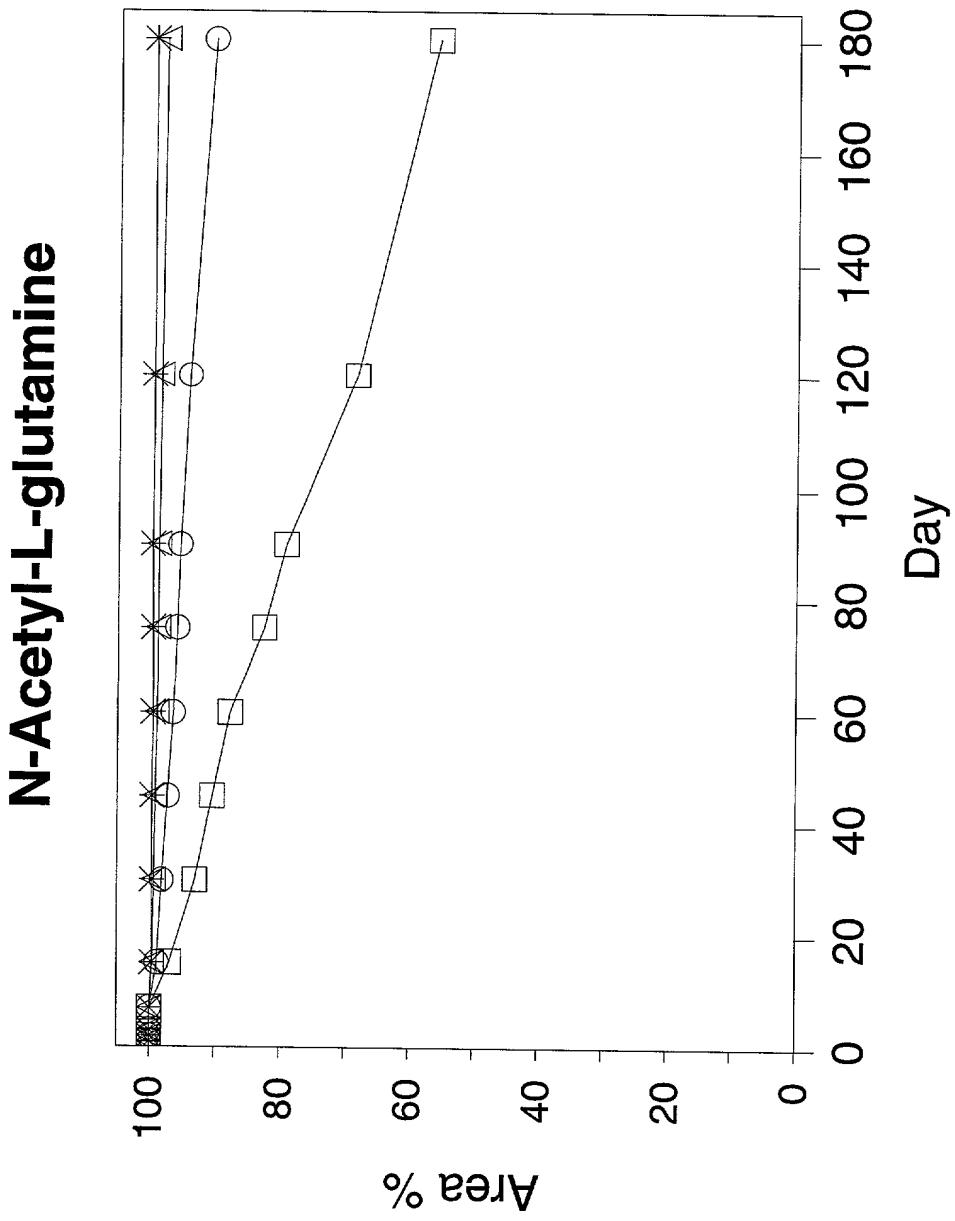


Figure 1: Aqueous stability of N-acetyl-L-glutamine at various pH (ambient temperature, in 1 pH unit increments from pH 2 to 8): (?) pH 2.0; (o) pH 3.0; (Δ) pH 4.0; (?) pH 5.0 to pH 8.0 (all values are the same for pH 5.0 to pH 8.0 samples).

Figure 2

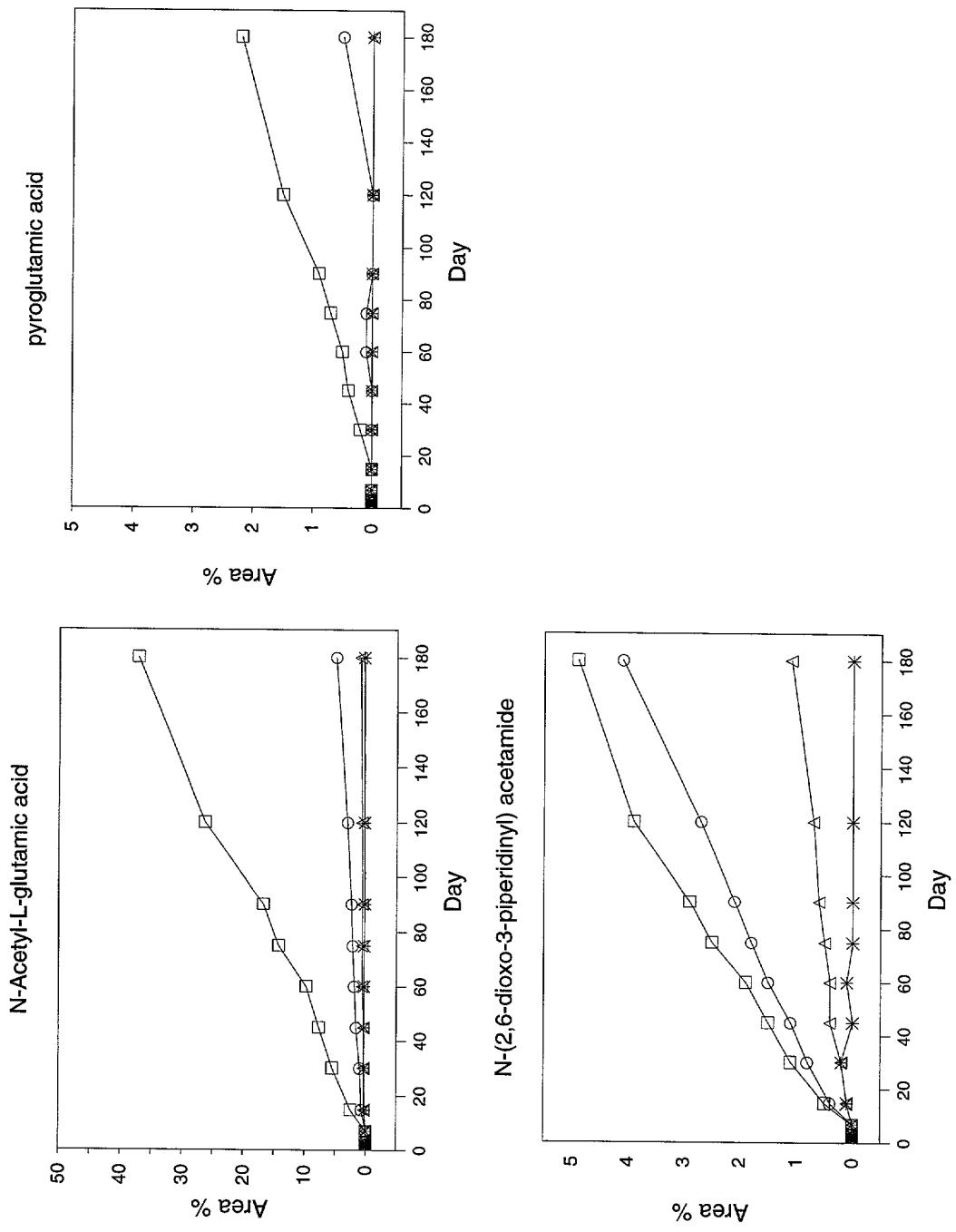
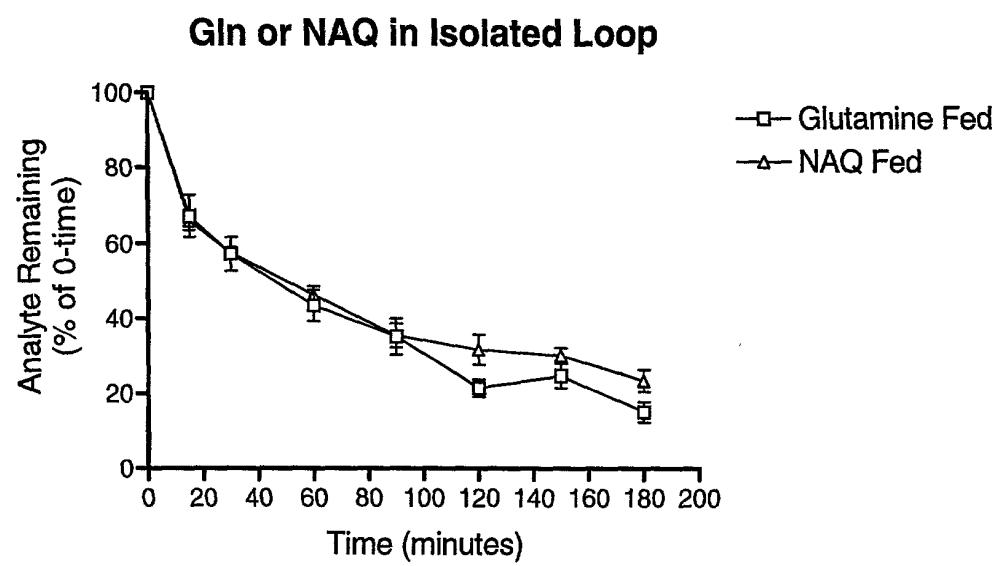
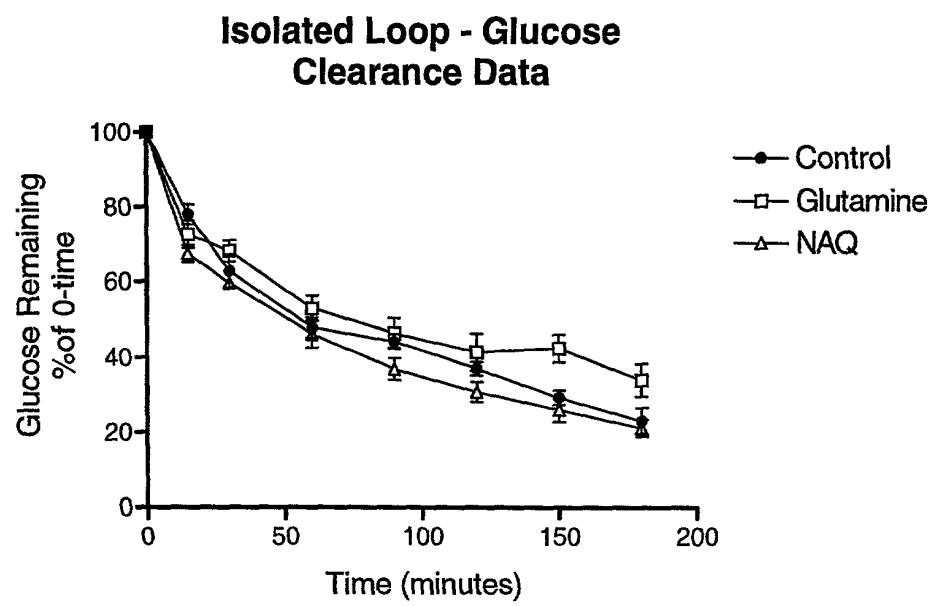


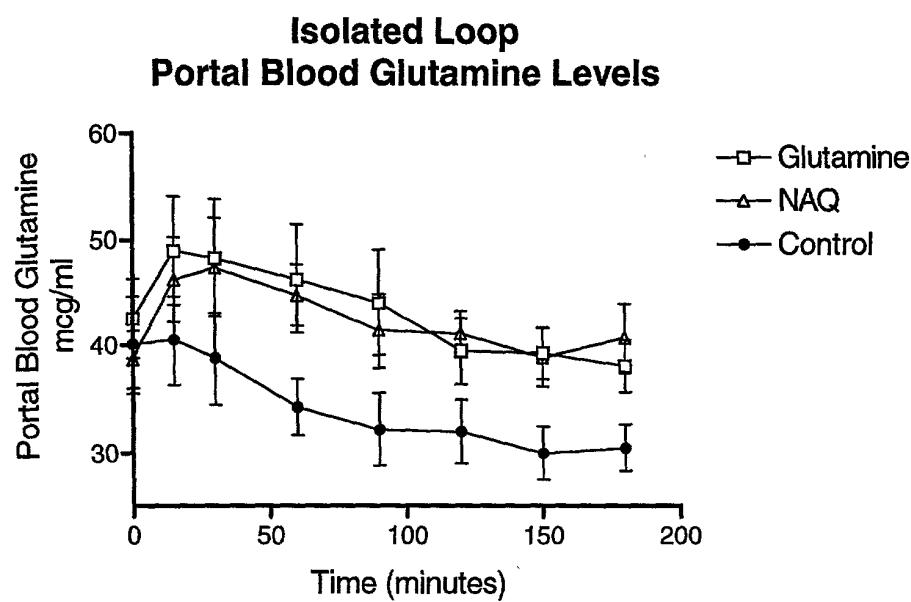
Figure 2: Degradation products of N-acetyl-L-glutamine in aqueous solution at various pH, (ambient temperature, in 1 pH unit increments from pH 2 to 8): (?) pH 2.0; (o) pH 3.0; (Δ) pH 4.0; (*) pH 5.0 to pH 8.0 (all values are the same for pH 5.0 to pH 8.0 samples).

Figure 3

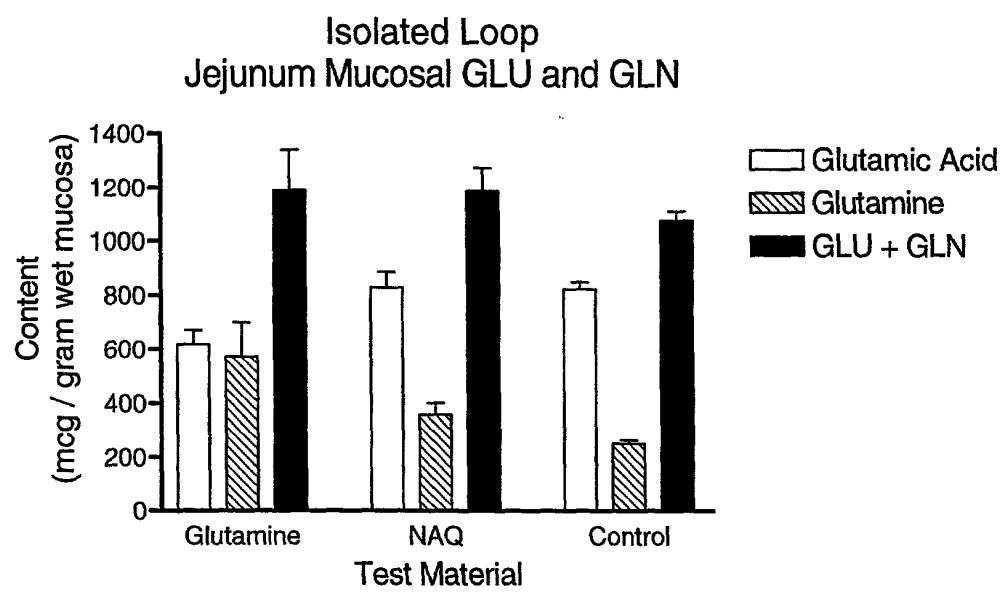
No significant difference between Gln or NAQ - $t_{1/2} \sim 45$ minutes.



Glucose clearance from isolated intestinal loop. There was no significant difference between groups, and $t_{1/2}$ is approximately 60 - 90 minutes.

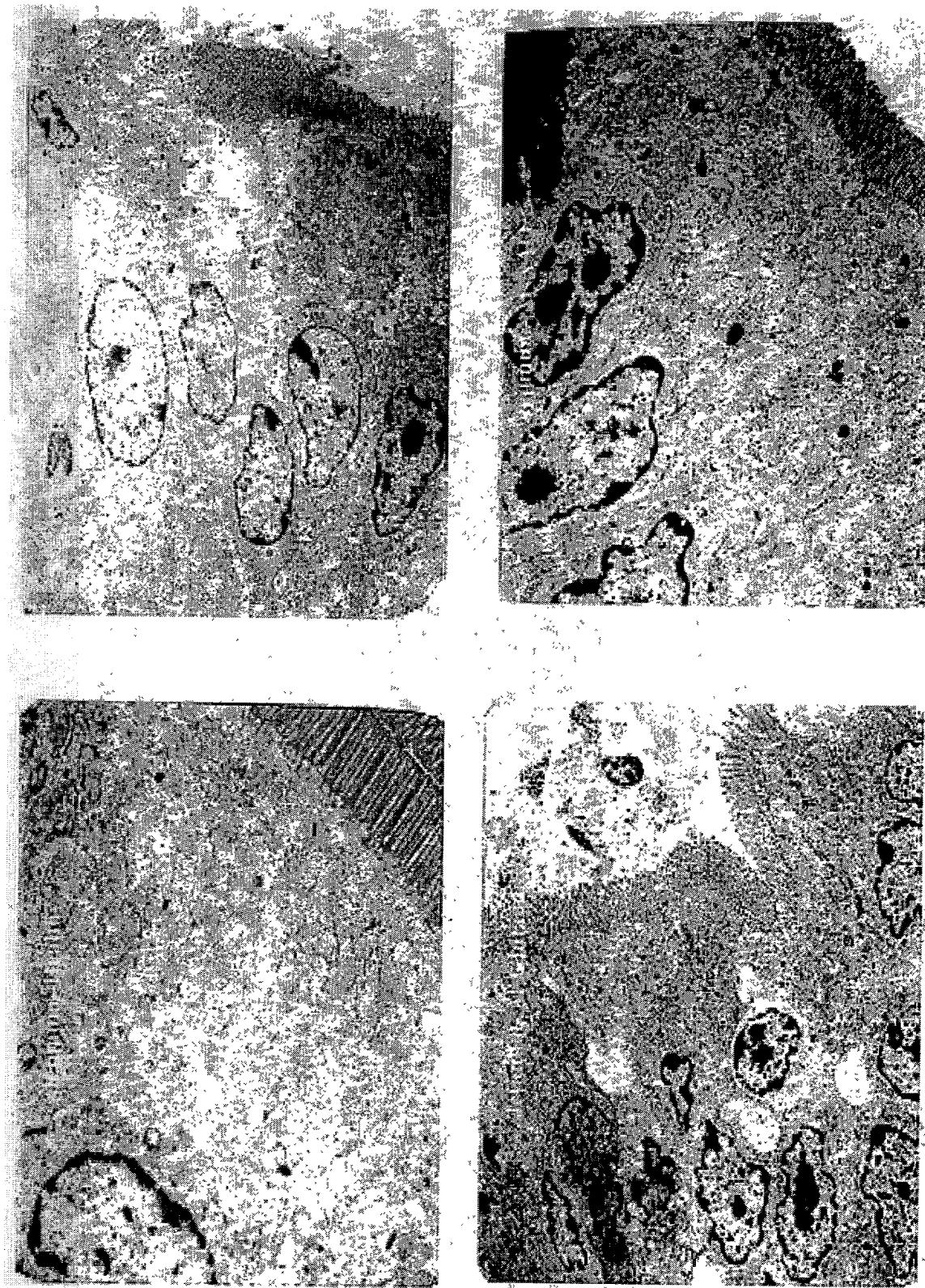


Portal blood glutamine levels are not different whether glutamine or NAQ is placed in the isolated intestinal loop. Both are different from the control (no added glutamine source).

Figure 6

Isolated Loop - Total mucosal glutamine and glutamate analysis.
 Incubation with NAQ or Gln results in similar retention of GLN+GLU in the mucosa. NAQ does not produce as high a GLN content in the mucosa as GLN, but NAQ does prevent the loss of GLU content observed with GLN. Both GLN and NAQ are identical in supporting a slightly elevated mucosal content of GLN+GLU compared to control.

Figure 7



Electron transmission micrographs of enterocyte cytoplasm from healthy and malnourished pigs. Malnourished pigs received daily a supplement of caseinate, glutamine or NAQ.